

**AMENDMENTS TO THE CLAIMS:**

This listing of claims will replace all prior versions and listings of claims in the application. Please cancel claims 1-8 and 10, and add new claims 14-18, as follows:

1-8. (Cancelled).

9. (Original). A method for prophylaxis of migraine which comprises administering a therapeutically effective amount of a selective dual antagonist for the 5-HT<sub>2B</sub> and 5-HT<sub>7</sub> receptors to a patient.

10. (Cancelled).

11. (Previously presented). The method of claim 9, wherein the selective dual antagonist for the 5-HT<sub>2B</sub> and 5-HT<sub>7</sub> receptors comprises:

a) a 5-HT<sub>2B</sub> receptor antagonistic compound as a first ingredient having a selective binding affinity to the 5-HT<sub>2B</sub> receptor, and

b) a 5-HT<sub>7</sub> receptor antagonistic compound as a second ingredient having a selective binding affinity to the 5-HT<sub>7</sub> receptor.

12. (Previously presented). The method of claim 9, wherein the selective dual antagonist for the 5-HT<sub>2B</sub> and 5-HT<sub>7</sub> receptors comprises a dual antagonistic compound for the 5-HT<sub>2B</sub> and 5-HT<sub>7</sub> receptors having a selective binding affinity to both of the 5-HT<sub>2B</sub> and 5-HT<sub>7</sub> receptors.

13. (Currently Amended). The method of claim 9, wherein the ~~binding affinity~~ Ki or IC<sub>50</sub> values for the 5-HT<sub>2B</sub> and 5-HT<sub>7</sub> receptors ~~[[is]]are~~ respectively one-hundredth or ~~more to the~~ less of those of each of  $\alpha_1$ , M<sub>1</sub>, D<sub>2</sub>, 5-HT<sub>1A</sub>, 5-HT<sub>1B</sub>, 5-HT<sub>2A</sub>, 5-HT<sub>2C</sub>, 5-HT<sub>3</sub>, 5-HT<sub>4</sub> and 5-HT<sub>6</sub> receptors.

14. (New). The method of claim 9, wherein the binding affinities for the 5-HT<sub>2B</sub> and 5-HT<sub>7</sub> receptors are higher than those of each of  $\alpha_1$ , M<sub>1</sub>, D<sub>2</sub>, 5-HT<sub>1A</sub>, 5-HT<sub>1B</sub>, 5-HT<sub>3</sub>, 5-HT<sub>4</sub> and 5-HT<sub>6</sub> receptors.

15. (New). The method of claim 9, wherein the binding affinities for the 5-HT<sub>2B</sub> and 5-HT<sub>7</sub> receptors are higher than those of each of  $\alpha_1$ , M<sub>1</sub>, D<sub>2</sub>, 5-HT<sub>1A</sub>, 5-HT<sub>1B</sub>, 5-HT<sub>2A</sub>, 5-HT<sub>2C</sub>, 5-HT<sub>3</sub>, 5-HT<sub>4</sub> and 5-HT<sub>6</sub> receptors.

16. (New). The method of claim 9, wherein the Ki or IC<sub>50</sub> values for the 5-HT<sub>2B</sub> and 5-HT<sub>7</sub> receptors are respectively one-tenth or less of those of each of  $\alpha_1$ , M<sub>1</sub>, D<sub>2</sub>, 5-HT<sub>1A</sub>, 5-HT<sub>1B</sub>, 5-HT<sub>3</sub>, 5-HT<sub>4</sub> and 5-HT<sub>6</sub> receptors.

17. (New). The method of claim 9, wherein the Ki or IC<sub>50</sub> values for the 5-HT<sub>2B</sub> and 5-HT<sub>7</sub> receptors are respectively one-tenth or less of those of each of  $\alpha_1$ , M<sub>1</sub>, D<sub>2</sub>, 5-HT<sub>1A</sub>, 5-HT<sub>1B</sub>, 5-HT<sub>2A</sub>, 5-HT<sub>2C</sub>, 5-HT<sub>3</sub>, 5-HT<sub>4</sub> and 5-HT<sub>6</sub> receptors.

18. (New). The method of claim 9, wherein the  $K_i$  or  $IC_{50}$  values for the 5-HT<sub>2B</sub> and 5-HT<sub>7</sub> receptors are respectively one-hundredth or less of those of each of  $\alpha_1$ , M<sub>1</sub>, D<sub>2</sub>, 5-HT<sub>1A</sub>, 5-HT<sub>1B</sub>, 5-HT<sub>3</sub>, 5-HT<sub>4</sub> and 5-HT<sub>6</sub> receptors.